

IDS Form PTO/SB/08: Substitute for form 1449A/PTO				Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)				Application Number	10/584,996
				Filing Date	June 29, 2006
				First Named Inventor	Robert DOBLHOFFER et al.
				Art Unit	
				Examiner Name	
Sheet	1	of	2	Attorney Docket Number	05281.0018

U.S. PATENTS AND PUBLISHED U.S. PATENT APPLICATIONS					
Examiner Initials	Cite No. ¹	Document Number Number-Kind Code ² (if known)	Issue or Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		US 5,922,713	07-13-1999	WERNER	

Note: Copies of the U.S. Patent Documents are not Required in IDS filed after October 21, 2004

FOREIGN PATENT DOCUMENTS						
Examiner Initials	Cite No. ¹	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	Translation ⁶
		Country Code ³ Number ⁴ Kind Code ⁵ (if known)				
		WO 95/32203	11-30-1995	PFLEIDERER et al.		Abstract
		DE 44 18 097 A1	11-30-1995	PFLEIDERER et al.		NO
		EP 0 906 913 A1	04-07-1999	WERNER et al.		NO
		WO 01/21619 A1	03-29-2001	PFLEIDERER et al.		Abstract
		WO 00/39129	07-06-2000	WAER et al.		
		GB 2 240 041 A1	07-24-1991	BRAQUET et al.		
		WO 93/13055	07-08-1993	BEAMS et al.		

NON PATENT LITERATURE DOCUMENTS					Translation ⁵
Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.			
		McCALL, T. B. et al., "Identification of N-Iminoethyl-L-Ornithine as an Irreversible Inhibitor of Nitric Oxide Synthase in Phagocytic Cells," Br. J. Pharmacol., Vol. 102, No. 1, p. 234, (1991). (Abstract Only)			
		MISKO, T. P. et al., "Selective Inhibition of the Inducible Nitric Oxide Synthase by Aminoguanidine," Eur. J. Pharmacol., Vol. 233, No. 1, p. 119, (1993). (Abstract Only)			
		MOORE, P. K. et al., "7-Nitro Indazole, an Inhibitor of Nitric Oxide Synthase, Exhibits Anti-Nociceptive Activity in the Mouse Without Increasing Blood Pressure," Br. J. Pharmacol., Vol. 106, No. 2, p. 296, (1993). (Abstract Only)			
		KWON, N. S. et al., "Reduced Biopterin as a Cofactor in the Generation of Nitrogen Oxides by Murine Macrophages," The Journal of Biological Chemistry, Vol. 264, No. 34, pp. 20496-20501, (1989).			
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		SCHIRCKS, V. B. et al., "A New, Regiospecific Synthesis of L-Bioterine," Helvetica Chimica Acta, Vol. 60, No. 1, pp. 211-214, (1977).		Abstract
		FUTTERMAN, S., "Enzymatic Reduction of Folic Acid and Dihydrofolic Acid to Tetrahydro-Folic Acid," J. Biol. Chem., Vol. 228, pp. 1031-1038, (1957). (Abstract Only)		
		FUKUSHIMA, T. et al., "Nuclear Magnetic Resonance Studies of Some Biologically Active Dihydropterins," Vol. 128, Issue 1, 1 page, (1968).		
		PFLEIDERER, W. et al., "A Simple Synthetic Approach to 8-Substituted 5,6,7,8-Tetrahydro- and 7,8-Dihydropterins," Chem. Ber., Vol. 104, pp. 2293-2312, (1971).		YES
		ANDREWS, K. J. M. et al., "A New Synthesis of Bioterine and L-Neopterine," Chemical Communications, pp. 120-121, (1968).		
		HANAYA, T. et al., "Pteridines CV Selective N(3)- and O'-Alkylation of L-Bioterine: A Convenient Synthesis of 3-and O'-Methyl-L-Bioterine and the Versatile N ⁶ -(N,N-Dimethylaminomethyl)-N(3)-p-Nitrophenethyl-Protected L-Bioterine," Pteridines, Vol. 6, No. 1, pp. 1-7, (1995).		
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		KWEE, S. et al., "Electrochemistry of Some Substituted Pteridines," Biochimica et Biophysica Acta, Vol. 297, No. 2, p 285-296, (1973). (Abstract Only)		
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Examiner Signature	/Cecilia Jaisle/	Date Considered	02/24/2008
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EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

ALL REFERENCES CONSIDERED EXCEPT WHERE LINED THROUGH. /CJ/